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<p>(54) Title: POLYSILICATE MICROGELS</p> <p>(57) Abstract</p> <p>The present invention generally relates to polysilicate microgels. More specifically, the invention relates to a process for preparing aqueous polysilicate microgels which comprises mixing an aqueous solution of alkali metal silicate with an aqueous phase of silica-based material having a pH of 11 or less, to polysilicate microgels per se and to the use thereof as flocculating agents in paper making and water purification. The invention further relates to a process for the production of paper from a suspension of cellulosic fibres, and optional filler, which comprises adding to the suspension at least one cationic or amphoteric organic polymer and a polysilicate microgel, forming and draining the suspension on a wire.</p>			

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### Polysilicate microgels

The present invention generally relates to polysilicate microgels. More particularly, the invention relates to polysilicate microgels, their preparation and use in paper making and water purification.

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#### Background

Polysilicate microgels, optionally aluminated, are known in the art as drainage and retention aids in the manufacture of paper and similar cellulosic products. The microgels comprise solutions or dispersions of very small primary silica-based particles, usually with a size of from 1 to 2 nm in diameter and having a high specific surface area, typically at least about 1000 m<sup>2</sup>/g, which are linked together into individual chains to form three-dimensional network structures.

The preparation of polysilicate and aluminated polysilicate microgels generally comprises acidifying a dilute aqueous solution of alkali metal silicate by using an acid or acid ion-exchanger, ageing the acidified mixture and then further diluting the aged mixture to a silica concentration of not greater than 2% by weight. The silica-based microgels normally have poor stability and the high dilution is normally necessary to avoid gelation of the microgels. Because of the stability problems associated with these products, and the prohibitive cost of shipping stable, but extremely dilute, solutions containing about 0.5% by weight or less of silica, the polysilicate microgels are preferably prepared at the location of intended use, for example at the paper mill. Production units or generators for continuously preparing polysilicate microgels that are installable at the paper mill are known in the art. Hereby the microgels obtained can be prepared and continuously introduced into the stock containing cellulosic fibres and filler to be drained. However, any disturbance in the production unit, for example variations in quality and/or quantity of the microgel produced, will change the drainage and retention performance of the product which may adversely affect the paper making process, thereby producing cellulosic products of uneven quality.

#### The Invention

30 In accordance with the present invention it has been found that polysilicate microgels can be prepared in a very advantageous manner by mixing an alkaline alkali metal silicate with a silica-based material. Hereby it is possible to prepare high-concentration polysilicate and aluminated polysilicate microgels and microgel precursors. More specifically, this invention concerns a process for preparing an aqueous polysilicate 35 microgel which comprises mixing an aqueous solution of alkali metal silicate with an aqueous phase of silica-based material preferably having a pH of 11 or less. The invention

thus relates to a process for preparing polysilicate microgels, polysilicate microgels per se and their use, as further defined in the claims.

The polysilicate microgels of this invention exhibit very high stability and can be easily prepared and shipped at considerably higher silica concentrations as compared to 5 previously known polysilicate microgels. High-concentration polysilicate microgels of this invention can be prepared under controlled conditions in a plant intended for such production and shipped as a concentrated product to the paper mill in an economically attractive manner. Hereby there will be no need for installing production units or generators for preparing polysilicate microgel drainage and retention aids in a large number of paper 10 mills, thereby offering substantial technical and economic benefits.

If desired, the high-concentration polysilicate microgel of this invention can be diluted with water or an aqueous acid solution before being used, for example as a drainage and retention aid in papermaking, in order to facilitate dosage of the silica-based material to the dilute fibre stock. In this respect, the high-concentration polysilicate microgel of the 15 invention can be regarded as a precursor for low-concentration polysilicate microgels which can be formed by the addition of water, optionally acidified. In this application, a storage tank for the high-concentration polysilicate microgel can be installed at the location of intended use, which is more attractive economically than installing a complete polysilicate microgel production unit or generator. Of course, low-concentration polysilicate 20 microgels can also be formed in situ in the aqueous phase into which the high-concentration silica-based product is being incorporated as a high-performance additive, for example by adding the high-concentration product to a cellulosic fibre suspension to be drained in papermaking or to waste water to be purified. This application using high-concentration polysilicate microgels of this invention as a precursor for in situ formation of 25 low-concentration polysilicate microgels represents a significant progress in the art.

The alkali metal silicate used to prepare the polysilicate microgels according to the invention can be any water-soluble silicate salt such as sodium or potassium silicate, or sodium or potassium water glass. These are available with varying molar ratios of  $\text{SiO}_2$  to 30  $\text{Na}_2\text{O}$  or  $\text{K}_2\text{O}$  and the molar ratio is usually within the range of from 1.5:1 to 4.5:1, most often from about 2.5:1 to 3.9:1. The alkali metal silicate preferably is a sodium silicate. Aqueous solutions of alkali metal silicate are alkaline and usually have a pH of about 13 or above 13. The alkali metal silicate solution usually have a silica concentration within the range of from 5 to 35% by weight, suitably above 10% by weight and preferably within the range of from 15 to 30% by weight.

35 The silica-based material to be mixed with the alkali metal silicate solution according to the invention can be selected from a wide variety of siliceous materials

including dispersed silicas such as, for example, silica-based sols, fumed silica, silica gels, precipitated silicas, acidified solutions of alkali metal silicates, and suspensions of silica-containing clays of smectite-type. The aqueous phase of silica-based material can have a pH within the range of from 1 to 11. In one preferred aspect of this invention, the pH of the aqueous silica-based material is within the range of from 1.5 to 4. In another preferred aspect of this invention, the pH of the aqueous silica-based material is within the range of from 4 to 11.0, usually from 4.5, suitably from 6.5 and most preferably from 7 up to 11.0, preferably up to 10.6.

In one preferred embodiment of this invention, the silica-based material used for mixing with the alkali metal silicate solution is a silica-based sol, suitably an alkali-stabilized silica sol. Sols of this type are known in the art and generally comprise an aqueous phase and particles based on silica, i.e.  $\text{SiO}_2$ . It is preferred that the particles are colloidal, i.e. in the colloidal range of particle size. Particles of this type include colloidal silica, colloidal aluminium-modified silica and colloidal aluminium silicate. The silica-based sols can have a pH as defined above and usually it is at least 4. Suitable sols with silica-based particles include those disclosed in U.S. Pat. Nos. 4,388,150; 4,961,825; 4,980,025; 5,368,833; 5,447,604; and 5,603,805; the teachings of which are incorporated herein by reference.

The silica-based particles contained in the sol can have an average particle size less than about 100 nm and suitably less than about 50 nm. In a preferred embodiment of the invention, the silica-based sol particles can have an average particle size of less than about 30 nm and preferably within the range of from about 1 to about 15 nm. As conventional in silica chemistry, the size refers to the average size of the primary particles, which may be aggregated or non-aggregated. The silica-based particles present in the sol should suitably have a specific surface area of at least  $50 \text{ m}^2/\text{g}$ . The specific surface area can be measured by means of titration with NaOH in known manner, e.g. as described by Sears in Analytical Chemistry 28(1956):12, 1981-1983 and in U.S. Pat. No. 5,176,891. The given area thus represents the average specific surface area of the particles. Suitably the specific surface area is within the range of from  $50$  to  $1200 \text{ m}^2/\text{g}$  and preferably from  $70$  to  $1000 \text{ m}^2/\text{g}$ .

In another preferred embodiment of this invention, the silica-based material used for mixing with the alkali metal silicate solution is an acidified alkali metal silicate. Suitable acidified alkali metal silicates include polysilicic acid, polymeric silicic acid, active, or activated, silica, and polysilicates, optionally aluminated. The acidified alkali metal silicate solution can be prepared in known manner by acidifying an aqueous solution of alkali metal silicate which may be any water-soluble silicate salt such as sodium or potassium silicate, or sodium or potassium water glass, preferably sodium silicate. Suitable alkali metal silicates are available with varying molar ratios of  $\text{SiO}_2$  to  $\text{Na}_2\text{O}$  or  $\text{K}_2\text{O}$  and the ratio is

usually within the range of from 1.5:1 to 4.5:1, most often from about 2.5:1 to 3.9:1. The acidified alkali metal silicate solution can have a pH as defined above. Acidification can be carried out in many ways, for example by using acid ion-exchange resins, mineral acids, e.g. sulphuric acid, hydrochloric acid or phosphoric acid, acid salts or acid gases, suitably ion-exchangers or mineral acids or a combination thereof, optionally in combination with an aluminium salt. Where higher ratios of SiO<sub>2</sub> to Na<sub>2</sub>O are desired, it is preferred to use acid ion-exchangers. Suitable acidified alkali metal silicates include those disclosed in U.S. Pat. Nos. 4,388,150; 4,954,220; 5,127,994; 5,279,807; 5,312,595; and 5,503,820; the teachings of which are incorporated herein by reference.

Suitable acidified alkali metal silicate solutions can also be selected from alkali metal silicates that have been both acidified and aluminated. The alumination can be carried out simultaneously with or after the acidification. Suitable aluminated and acidified alkali metal silicates include aluminated polysilicates or polyaluminosilicates, for example as disclosed in U.S. Pat. Nos. 5,176,891; 5,470,435; 5,482,693; 5,543,014; and 5,626,921, the teachings of which are incorporated herein by reference.

Acidified, and optionally aluminated, alkali metal silicates for use in the process of this invention suitably have a high specific surface area, suitably above about 1000 m<sup>2</sup>/g. The specific surface area can be within the range of from 1000 to 1700 m<sup>2</sup>/g, preferably from 1050 to 1600 m<sup>2</sup>/g. The specific surface area can be measured as described above.

The process of this invention comprises mixing the alkaline alkali metal silicate with the silica-based material. The mixing can be conducted at a temperature of from 0 to 300°C, suitably from 5 to 150°C and preferably from about 10 to about 100°C. The process can be carried out by adding the silica-based material to the aqueous solution of alkali metal silicate with stirring. Preferably the silica-based material is slowly added to the alkali metal silicate under vigorous mixing. The aqueous phase of silica-based material can have a silica concentration within the range of from 0.1 to 60% by weight, depending on the type of material used. When using silica-based sols such as alkali-stabilized sols, the SiO<sub>2</sub> content can be from 5 to 60% by weight, preferably from 10 to 50% by weight. When using acidified alkali metal silicates as described herein, the SiO<sub>2</sub> content can be from 0.1 to 10% by weight, preferably from 0.5 to 6% by weight. The SiO<sub>2</sub> present in the polysilicate microgels obtained thus originates from both the alkali metal silicate and the silica-based material. The mixing can be carried out such that from 5 to 95% by weight and suitably from 15 to 85% by weight of SiO<sub>2</sub> present in the polysilicate microgel is derived from the alkali metal silicate, and 95 to 5% by weight and suitable from 85 to 15% of SiO<sub>2</sub> present in the polysilicate microgel is derived from the silica-based material.

If desired, additional compounds can be incorporated into the mixture obtained in the process. Suitable additional compounds include various salts such as, for example, aluminium salts and other metal salts, and various acids. These additional compounds may have a positive effect on storage stability and/or on the drainage and/or retention 5 performance of the microgels. Suitable aluminium salts include alum, aluminates, aluminium chloride, aluminium nitrate and polyaluminium compounds, such as polyaluminium chlorides, polyaluminium sulphates, polyaluminium compounds containing both chloride and sulphate ions, polyaluminium silicate-sulphates, and mixtures thereof. The polyaluminium compounds may also contain other anions, for example anions from phosphoric acid, organic 10 acids such as citric acid and oxalic acid. Preferred aluminium salts include aluminates, e.g. sodium or potassium aluminate, preferably sodium aluminate. Aluminium salts can be incorporated into the polysilicate microgels in an amount corresponding to a  $\text{SiO}_2:\text{Al}_2\text{O}_3$  molar ratio within the range of from 1500:1 to 5:1 and suitably from 750:1 to 6:1. Suitable metal salts other than salts based on Al include those based on alkali metals and alkaline earth 15 metals such as, for example, Li, K, Mg and Ca. Suitable anions can be selected from hydroxide, borate, nitrate, chloride, formate, acetate, etc. These additional compounds are generally incorporated in amounts such that the ratio of Na to such additional metal present in the metal salt is higher than 1, preferably higher than 5. Suitable acids include organic acids, preferably diacids such as dicarboxylic and disulphonic acids, e.g. oxalic, malonic, 20 succinic, glutaric and adipic acid, and organic polyacids, such as polymers containing carboxylic acid and sulphonic acid groups, such as polyacrylic acid. The additional acid is suitably incorporated in the polysilicate microgels in an amount sufficient to give a positive effect on stability, suitably up to 10% by weight. Even if arbitrary order of mixing or addition can be used, it is generally preferred that these additional compound be incorporated in the 25 mixture of alkali metal silicate and silica-based material. A further preferred order of mixing is to add the acid to the acidified solution of alkali metal silicate prior to mixing with the alkali metal silicate solution.

The resulting aqueous solution, or dispersion, of polysilicate microgel, or microparticulate silica-based material, optionally aluminated, normally have a  $\text{SiO}_2$  content 30 of at least 5% by weight. The  $\text{SiO}_2$  content may depend on several factors such as, for example, the starting materials used and their  $\text{SiO}_2$  content, the incorporation of aluminium salts, etc. Suitably the  $\text{SiO}_2$  content is at least 10% by weight, preferably at least 15% by weight and most preferably at least 17.5% by weight. The upper limit is usually about 50% by weight, suitably about 35% by weight and in most cases about 30% by weight. If 35 desired, after preparation, the aqueous polysilicate microgel can be subjected to further treatment like ion-exchange and/or concentration. This can be advantageous so as to

provide products having improved storage-stability and/or higher concentration. Concentration can be carried out by known methods, for example by membrane processes or evaporation of water.

The polysilicate microgel obtained can have a molar ratio of  $\text{SiO}_2:\text{Na}_2\text{O}$  within the range of from 3:1, suitably from 4:1, and preferably from 5:1; up to 50:1, suitably to 30:1 and preferably to 20:1. This ratio may depend on several factors such as, for example, type of starting materials, type of acidification of starting materials, additional metal salts, etc. The same applies the molar ratio of  $\text{SiO}_2:\text{M}_2\text{O}$ , where M is alkali metal, e.g. Li, Na, K and mixtures thereof. The polysilicate microgel can have a molar ratio of  $\text{SiO}_2:\text{M}_2\text{O}$  within the range of from 2:1, suitably from 2.2:1, and preferably from 3:1, up to 50:1, suitably to 30:1, and preferably to 20:1. Preferably the polysilicate microgel is anionic in nature. The obtained aqueous polysilicate microgel, or silica-based microparticulate material, generally have a pH below 14, suitably below 13 and preferably below 12. Usually pH is above 6 and suitably above 9.

The high-concentration polysilicate microgel of this invention can be diluted or mixed with aqueous solutions or suspensions before use or at use. Hereby low-concentration polysilicate microgels can be formed and the high-concentration polysilicate microgel of the invention can thus be used as a precursor for low-concentration polysilicate microgels. In a preferred embodiment, the high-concentration polysilicate microgel is mixed with an aqueous solution or suspension having a pH lower than that of the high-concentration microgel whereby the high-concentration microgel undergoes a pH transition. Suitable aqueous solutions and suspensions include those containing acids, cellulosic fibres and optional filler, and acid aluminium salts.

The polysilicate microgel obtained by the process can be described as a silica-based microparticulate material comprising very small particles, preferably 1-2 nm in diameter, which are linked together in chains or networks to form three-dimensional structures. The aqueous polysilicate microgels may also contain larger particles depending on, inter alia, the starting materials used in the preparation of the microgels. The specific surface area of the silica-based microparticles, the microgel, is suitably at least 1000  $\text{m}^2/\text{g}$  and usually up to about 1700  $\text{m}^2/\text{g}$ . Methods for measuring the specific surface area are described hereinabove.

The polysilicate microgels of this invention are suitable for use as flocculating agents, for example in the production of pulp and paper and within the field of water purification, both for purification of different kinds of waste water and for purification specifically of white water from the pulp and paper industry. The polysilicate microgels can be used as flocculating agents in combination with organic polymers which can be selected from

anionic, amphoteric, nonionic and cationic polymers and mixtures thereof. The use of such polymers as flocculating agents is well known in the art. The polymers can be derived from natural or synthetic sources, and they can be linear or branched. Examples of generally suitable polymers include anionic, amphoteric and cationic starches, anionic, amphoteric and 5 cationic guar gums, and anionic, amphoteric and cationic acrylamide-based polymers, as well as cationic poly(diallyldimethyl ammonium chloride), cationic polyethylene imines, cationic polyamines, polyamidoamines and vinylamide-based polymers, melamine-formaldehyde and urea-formaldehyde resins. Suitably the polysilicate microgel is used in combination with at least one cationic or amphoteric polymer, preferably cationic polymer. Cationic starch and 10 cationic polyacrylamide are particularly preferred polymers and they can be used singly, together with each other or together with other polymers, e.g. other cationic polymers or anionic polyacrylamide. Even if arbitrary order of addition can be used, it is preferred that the polymer or polymers be added to pulp, stock or water before the polysilicate microgels.

The preferred field of use for the polysilicate microgels, in combination with polymer 15 as described above, is for improvement of drainage and/or retention in the manufacture of paper, i.e. the use as drainage and/or retention aids in papermaking. The present invention further relates to a process for the production of paper from a suspension of cellulosic fibres, and optional fillers, which comprises adding to the suspension at least one cationic or amphoteric organic polymer and polysilicate microgel as described herein, forming and 20 draining the suspension on a wire. The invention thus relates to a process as further defined in the claims.

When using the polysilicate microgels in combination with organic polymer(s) as mentioned above, it is further preferred to use at least one anionic trash catcher (ATC). ATC's are known in the art as neutralizing agents for detrimental anionic substances present 25 in the stock. Hereby ATC's can enhance the efficiency of other additives used in the process. Accordingly, further suitable combinations of polymers that can be co-used with the polysilicate microgels of this invention include ATC in combination with high molecular weight polymer(s), e.g. cationic starch and/or cationic polyacrylamide, anionic polyacrylamide as well as cationic starch and/or cationic polyacrylamide in combination with anionic polyacrylamide. 30 Suitable ATC's include cationic polyelectrolytes, especially low molecular weight highly charged cationic organic polymers such as polyamines, polyethyleneimines, homo- and copolymers based on diallyldimethyl ammonium chloride, (meth)acrylamides and (meth)-acrylates. Normally, ATC's are added to the stock prior to other polymer(s). Alternatively, the ATC polymer can be added simultaneously with the other polymer(s), either separately or in 35 admixture, for example as disclosed in European Pat. Appl. No. 752496, the teaching of

which is incorporated herein by reference. Mixtures comprising ATC polymer and high molecular weight cationic polymer are particularly preferred.

The amount of polysilicate microgel added to the stock, or suspension of cellulosic fibres, may vary within wide limits depending on, among other things, type of stock, type of polysilicate microgel used. The amount usually is at least 0.01 kg/ton and often at least 0.05 kg/ton, calculated as  $\text{SiO}_2$  and based on dry stock system, i.e. cellulosic fibres and optional fillers. The upper limit can be 8 kg/ton and suitably is 5 kg/ton. Usually the polysilicate microgel dosage is within the range of from 0.1 to 2 kg/ton.

The dosage of organic polymer to the stock can be varied over a broad range depending on, among other things, the type of polymer or polymers used and whether other effects are desired, e.g. wet and dry paper strength. Usually, there is used at least 0.005 kg of polymer per ton of dry fibres and optional fillers. For synthetic cationic polymers, such as for example cationic polyacrylamide, amounts of at least 0.005 kg/ton are usually used, calculated as dry on dry fibres and optional fillers, suitably from 0.01 to 3 and preferably from 0.03 to 2 kg/ton. For cationic polymers based on carbohydrates, such as cationic starch and cationic guar gum, amounts of at least 0.05 kg/ton, calculated as dry on dry fibres and optional fillers, are usually used. For these polymers the amounts are suitably from 0.1 to 30 kg/ton and preferably from 1 to 15 kg/ton.

In a preferred embodiment of this invention, the polysilicate microgels are used as drainage and/or retention aids in combination with at least one organic polymer, as described above, and at least one aluminium compound. Aluminium compounds can be used to further improve the drainage and/or retention performance of stock additives comprising polysilicate microgels and aluminated polysilicate microgels. Suitable aluminium salts include alum, aluminates, aluminium chloride, aluminium nitrate and polyaluminium compounds, such as polyaluminium chlorides, polyaluminium sulphates, polyaluminium compounds containing both chloride and sulphate ions, polyaluminium silicate-sulphates, and mixtures thereof. The polyaluminium compounds may also contain other anions, for example anions from phosphoric acid, organic acids such as citric acid and oxalic acid. Preferred aluminium salts include sodium aluminate, alum and polyaluminium compounds. The aluminium compound can be added before, simultaneously with or after the addition of the polysilicate microgel. In many cases, it is often suitable to add the aluminium compound to the stock early in the process, for example prior to the other additives. Alternatively, or additionally, the aluminium compound can be added simultaneously with the polysilicate microgel at essentially the same point, either separately or in admixture with it, for example as disclosed in European Pat. Appl. No. 748897, the teachings of which is incorporated herein by reference. Adding the

polysilicate microgels and aluminium compound simultaneously to the stock represent a preferred embodiment of the invention.

The amount of aluminium compound added to the suspension may depend on the type of aluminium compound used and whether other effects are desired. It is for instance 5 well-known in the art to utilize aluminium compounds as precipitants for rosin-based sizes. The amount of aluminium compound added to the stock should suitably be at least 0.001 kg/ton, calculated as  $\text{Al}_2\text{O}_3$  and based on dry fibres and optional fillers. Suitably the amount is within the range of from 0.01 to 5 kg/ton and preferably from 0.05 to 1 kg/ton.

The papermaking process according to the invention can be used for producing 10 cellulosic products in sheet or web form such as for example pulp sheets and paper. It is preferred that the process is used for the production of paper. The term "paper", as used herein, of course include not only paper and the production thereof, but also other sheet or web-like products, such as for example board and paperboard, and the production thereof. The papermaking process according to the invention can be used in the production of sheet 15 or web-like products from different types of suspensions containing cellulosic fibres and the suspension, or stock, should suitably contain at least 50% by weight of such fibres, based on dry substance. The suspension can be based on fibres from chemical pulp, such as sulphate and sulphite pulp, thermomechanical pulp, chemo-thermomechanical pulp, refiner pulp or groundwood pulp from both hardwood and softwood, and can also be used for suspensions 20 based on recycled fibres. The suspension can also contain mineral fillers of conventional types, such as for example kaolin, titanium dioxide, gypsum, talc and both natural and synthetic calcium carbonates. The suspension can have a pH within the range from about 3 to about 10. The pH is suitably above 3.5 and preferably within the range of from 4 to 9. The stock can of course also contain papermaking additives of conventional types, such as wet- 25 strength agents, stock sizes based on rosin, ketene dimers or alkenyl succinic anhydrides, and the like.

The invention is further illustrated in the following Examples which, however, are not intended to limit same. Parts and % relate to parts by weight and % by weight, respectively, unless otherwise stated.

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#### Example 1

Polysilicate microgels of this invention were prepared by slowly adding a silica-based sol to sodium water glass (hereafter Na silicate) having a  $\text{SiO}_2$  content of about 28% by weight and a molar ratio of  $\text{SiO}_2:\text{Na}_2\text{O}$  of about 3.3 at 50°C under vigorous stirring. In 35 some tests, water and/or an additional alkali metal salt, either potassium borate ( $\text{KBO}_2$  (aq);

57% active) and/or lithium hydroxide (LiOH (aq); 97% LiOH·H<sub>2</sub>O), were introduced into the mixture thus obtained. The silica-based sols used were the following:

A: A sol of colloidal silica of the type described in U.S. Pat. No. 4,388,150. The sol was alkali-stabilized to a molar ratio of SiO<sub>2</sub>:Na<sub>2</sub>O of about 40, had a pH of about 8-9 and a SiO<sub>2</sub> content of about 15% by weight and contained silica particles with a specific surface area of about 500 m<sup>2</sup>/g.

B: A sol of colloidal silica having a SiO<sub>2</sub> content of 40% by weight and containing silica particles with a specific surface area of 220 m<sup>2</sup>/g. This sol is commercially available under the tradename Bindzil™ 40/220, Eka Chemicals.

10 C: A sol of aluminium-modified silicic acid of the type described in U.S. Pat. Nos. 4,961,825 and 4,980,025. The sol had a pH of about 8-9 and a SiO<sub>2</sub> content of about 15% by weight and contained silica particles with a specific surface area of about 500 m<sup>2</sup>/g.

15 Table I below shows the amounts (Amt.(g)) of starting materials used in the process, and the molar ratios of SiO<sub>2</sub>:Na<sub>2</sub>O and SiO<sub>2</sub>:M<sub>2</sub>O (SiO<sub>2</sub>:Na<sub>2</sub>O/SiO<sub>2</sub>:M<sub>2</sub>O), where M is alkali metal (Na+K+Li), and the content of SiO<sub>2</sub>, in % by weight (SiO<sub>2</sub> (%)), of the aqueous polysilicate microgel obtained.

Table I

Product No.	Na silicate Amt.(g)	SiO <sub>2</sub> sol Type/Amt.(g)	KBO <sub>2</sub> (aq) Amt (g)	LiOH (aq) Amt.(g)	SiO <sub>2</sub> :Na <sub>2</sub> O/	SiO <sub>2</sub>
					SiO <sub>2</sub> :M <sub>2</sub> O	(%)
20	1	300	A/319	18.8	5.0/3.4	19.5
	2	300	A/319	-	5.0/3.8	19.5
	3	500	A/150	-	3.8/3.8	24.1
	4	500	B/180	30.4	5.0/3.4	23.0
	5	500	B/180	-	5.0/3.8	23.0
25	6	400	B/231	-	6.0/4.3	23.0
	7	300	C/325	-	5.0/3.8	19.4
	8	300	C/325	-	5.0/3.4	16.1
	9	300	C/325	-	5.0/3.8	19.0
	10	500	C/153	-	3.8/3.8	19.6

30

Example 2

35 Stability of polysilicate microgels according to Example 1 was evaluated by measuring the viscosity 1, 3, 5 or 10 days after preparation. The viscosity measurements were made at 20°C with a Brookfield viscosimeter model RTV, spindle 61, 60 rpm. The results are set forth in Table II.

Table II

Product No.	SiO <sub>2</sub> (%)	Viscosity (cP) at 20°C			
		1 day	3 days	5 days	10 days
5	1	19.5	-	14	14
	2	19.5	-	11	11
	3	24.1	-	55	58
	4	23.0	29	-	52
	5	23.0	19	-	28
10	6	23.0	27	-	34
	7	19.4	11	-	6
	8	16.1	6	-	6
	9	19.0	7	-	7
	10	19.6	12	-	12

**15 Example 3**

In the following tests, drainage and retention performance of polysilicate microgels according to Example 1 was tested. Drainage performance was evaluated by means of a Dynamic Drainage Analyser (DDA), available from Akribi, Sweden, which measures the time for draining a set volume of stock through a wire when removing a plug and applying a vacuum to that side of the wire opposite to the side on which the stock is present. Retention performance was evaluated by means of a nephelometer by measuring the turbidity of the filtrate, the white water, obtained by draining the stock.

The tests were made using an alkaline stock at pH 8.0 and 0.25% consistency containing cellulosic fibres (60% bleached birch/40% pine sulphate) and 30% of chalk to which stock 0.3 g/l of Na<sub>2</sub>SO<sub>4</sub>·10H<sub>2</sub>O was added. In the tests, the polysilicate microgels were tested in conjunction with a cationic polymer, Raisamyl 142, which is a conventional medium-high cationized starch having a degree of substitution of 0.042, which was added to the stock in an amount of 12 kg/ton, calculated as dry on dry stock system, and an aluminium salt, sodium aluminate, which was added in an amount of 0.2 kg/ton, calculated as Al<sub>2</sub>O<sub>3</sub> and based on dry stock system.

The stock was stirred in a baffled jar at a speed of 1500 rpm throughout the test and chemical additions to the stock were conducted as follows:

- i) adding cationic starch followed by stirring for 30 seconds,
- ii) adding polysilicate microgels and aluminium salt simultaneously but separately followed by stirring for 15 seconds,
- iii) draining the stock while automatically recording the drainage time.

Table III shows the results obtained when using varying dosages (kg/ton, calculated as SiO<sub>2</sub> and based on dry stock system) of polysilicate microgels. Without addition of chemicals, the stock had a drainage time of 19.5 sec and a turbidity of 93 NTU.

Table III

5	Product	Drainage time (sec) / Turbidity (NTU) at SiO <sub>2</sub> dosage of				
		No.	0.5 kg/ton	0.8 kg/ton	1.2 kg/ton	1.7 kg/ton
	1		15.0/-	10.5/53	8.7/56	7.2/48
	2		16.5/-	12.3/73	9.9/61	9.1/62
	3		15.2/-	14.5/-	12.8/-	12.0/-
10	4		14.3/-	10.9/69	8.0/55	6.5/52
	6		14.6/-	10.3/65	8.2/60	7.5/55
	8		17.2/-	15.0/-	13.1/-	11.5/-
						10.7/-

Example 4

Polysilicate microgels of this invention were prepared by mixing at room temperature sodium water glass (hereafter Na silicate) having a SiO<sub>2</sub> content of about 23% by weight and a molar ratio SiO<sub>2</sub>:Na<sub>2</sub>O of about 3.3 with an acid ion-exchanger treated sodium silicate solution (Acid SiO<sub>2</sub>) having a SiO<sub>2</sub> content of about 5-6% by weight and pH of about 2.5 to achieve a mixture with a molar ratio SiO<sub>2</sub>:Na<sub>2</sub>O of about 4.0, followed by slowly adding a sodium aluminate solution (NaAl (aq)) containing about 2.5% by weight of Al<sub>2</sub>O<sub>3</sub> under vigorous stirring, or by adding an aqueous solution of an additional alkaline earth salt, either magnesium acetate (M; 1.5% Mg(acetate)<sub>2</sub>) or calcium nitrate (C; 2.5% Ca(NO<sub>3</sub>)<sub>2</sub>).

Table IV shows the amounts (Amt.(g)) of the starting materials used in the process, and the pH and silica content, in % by weight, of the polysilicate microgels obtained.

Table IV

25	Product	Na silicate	Acid SiO <sub>2</sub>	NaAl (aq.)	Salt (aq)	pH	SiO <sub>2</sub>
		No.	Amt. (g)	Amt. (g)	Amt (g)	Type/Amt.(g)	(%)
	11	320	237	34.3	-	11.3	18.8
	12	140	250	-	M/57	10.9	12.1
	13	129	100	-	C/21	11.5	13.9

Example 5

The polysilicate microgels according to Example 5 were tested for drainage and retention performance as in Example 3 using a similar stock of pH 8.4 and the same cationic starch, aluminium salt, dosages and order of addition.

Table V shows the results obtained by using polysilicate microgels in varying amounts, in kg/ton, calculated as SiO<sub>2</sub> and based on dry stock system. Without addition of chemicals, the stock had a drainage time of 21.2 sec and a turbidity of 104 NTU.

Table V

Product No.	Drainage time (sec) / Turbidity (NTU) at SiO <sub>2</sub> dosage of			
	0.5 kg/ton	0.8 kg/ton	1.2 kg/ton	1.7 kg/ton
5 11	15.2/84	12.0/74	11.6/66	7.9/ -
	12	13.9/82	11.1/70	9.9/62
	13	15.7/81	11.1/72	9.7/64

Example 6

Aluminated polysilicate microgels according to the invention were prepared by mixing sodium water glass with a SiO<sub>2</sub> content of about 23% by weight and molar ratio SiO<sub>2</sub>:Na<sub>2</sub>O of about 3.3 with an acid ion-exchanger treated sodium silicate solution having a SiO<sub>2</sub> content of about 5-6% by weight and pH of about 2.5 to achieve a mixture having a molar ratio SiO<sub>2</sub>:Na<sub>2</sub>O of about 10:1 followed by adding a dilute sodium aluminate solution (NaAl (aq)) containing about 2.5% by weight of Al<sub>2</sub>O<sub>3</sub> under stirring.

Table VI shows the SiO<sub>2</sub> and Al<sub>2</sub>O<sub>3</sub> contents of the products obtained as well as their stability and molar ratio of SiO<sub>2</sub>:Na<sub>2</sub>O. Stability was evaluated visually about 1 week after preparation.

Table VI

Product No.	SiO <sub>2</sub> (%)	Al <sub>2</sub> O <sub>3</sub> (%)	Stability	SiO <sub>2</sub> :Na <sub>2</sub> O (molar ratio)
20 14	10.7	0.4	OK	7.5
15	10.7	0.6	OK	6.2

Example 7

The polysilicate microgels according to Example 6 were tested for drainage and retention performance as in Example 3 using a similar stock at pH 8.4 and 0.27% consistency, except that no aluminium salt was added separately to the stock. The cationic starch and order of addition used in Example 3 were similarly used in this test series.

Table VII shows the results obtained when adding the polysilicate microgels in varying amounts, in kg/ton, calculated as SiO<sub>2</sub> and based on dry stock system. Without addition of chemicals, the stock had a drainage time of 21.5 sec and a turbidity of 97 NTU.

Table VII

Product No.	Drainage time (sec) / Turbidity (NTU) at SiO <sub>2</sub> dosage of				
	0.3 kg/ton	0.6 kg/ton	1.0 kg/ton	1.5 kg/ton	2.0 kg/ton
14	16.5/81	12.0/83	9.0/56	8.0/47	7.5/ -
15	16.0/72	12.2/ -	9.0/54	7.5/49	7.5/44

Claims

1. A process for preparing an aqueous polysilicate microgel, characterised in that it comprises mixing an aqueous solution of alkali metal silicate with an aqueous phase of silica-based material having a pH of 11 or less.
- 5 2. A process according to claim 1, characterised in that the aqueous polysilicate microgel obtained has a SiO<sub>2</sub> content of at least 15%.
3. A process according to claim 1 or 2, characterised in that the aqueous polysilicate microgel obtained comprises silica-based particles with a size of about 1 to 2 nm in diameter which are linked together in chains or networks to form three-dimensional
- 10 10. structures.
4. A process according to claim 1, 2 or 3, characterised in that the polysilicate microgel has a specific surface area of at least 1000 m<sup>2</sup>/g.
5. A process according to claim 1, 2, 3 or 4, characterised in that the aqueous silica-based material has a pH within the range of from 4 to 11.
- 15 6. A process according to claim 5, characterised in that the aqueous silica-based material has a pH within the range of from 7 to 11.
7. A process according to any of the preceding claims, characterised in that the aqueous silica-based material is a silica-based sol.
- 20 8. A process according to any of the preceding claims, characterised in that the silica-based material contains aluminium.
9. A process according to any of the preceding claims, characterised in that it further comprises admixing an additional salt.
10. A process according to claim 9, characterised in that the salt is an aluminium salt.
- 25 11. A process according to claim 9 or 10, characterised in that the salt is a metal salt other than an aluminium salt and based on an alkali metal or alkaline earth metal.
12. A process according to any of the preceding claims, characterised in that the polysilicate microgel obtained has a molar ratio SiO<sub>2</sub>:M<sub>2</sub>O, where M is alkali metal,
- 30 30. between 3:1 and 20:1.
13. Polysilicate microgel obtainable by a process according to any one of claims 1 to 12.
14. Use of a polysilicate microgel according to claim 13 or prepared by a process according to any of claims 1 to 12 as a flocculating agent in combination with at least one cationic or amphoteric polymer in the production of pulp and paper and for water
- 35 35. purification.

15. Use according to claim 14, wherein the polysilicate microgel is used as flocculating agent for improving drainage and/or retention in paper making.

16. Use according to claim 14 or 15, wherein the polysilicate microgel is used in combination with cationic starch and/or cationic acrylamide-based polymer.

5        17. A process for the production of paper from a suspension of cellulosic fibres, and optional filler, **c h a r a c t e r i s e d** in that comprises adding to the suspension at least one cationic or amphoteric organic polymer and a silica-based microparticulate material, forming and draining the suspension on a wire, **c h a r a c t e r i s e d** in that the silica-based material is a polysilicate microgel according to claim 13 or prepared by a process according  
10 to any of claims 1 to 12.

18. A process according to claim 17, **c h a r a c t e r i s e d** in that it further comprises adding an aluminium salt to the suspension.

# INTERNATIONAL SEARCH REPORT

International Application No  
PCT/SE 98/01101

**A. CLASSIFICATION OF SUBJECT MATTER**  
IPC 6 C01B33/143 D21H21/10 D21H17/68 C02F1/52

According to International Patent Classification(IPC) or to both national classification and IPC

**B. FIELDS SEARCHED**

Minimum documentation searched (classification system followed by classification symbols)  
IPC 6 C01B

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

**C. DOCUMENTS CONSIDERED TO BE RELEVANT**

Category °	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	<p>WO 94 05596 A (EKA NOBEL AB) 17 March 1994</p> <p>see claims 1,3-10 see page 1, line 3 - line 10 see page 2, line 24 - page 3, line 29 see page 4, line 6 - page 7, line 19 see page 8, line 10 - line 19 see example 1</p> <p>-----</p> <p>&amp; US 5 603 805 A cited in the application</p> <p>-----</p> <p>-----</p>	1,5, 7-10, 13-18
A		2-4,6,12

Further documents are listed in the continuation of box C.

Patent family members are listed in annex.

° Special categories of cited documents :

- "A" document defining the general state of the art which is not considered to be of particular relevance
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Name and mailing address of the ISA

European Patent Office, P.B. 5818 Patentlaan 2  
NL - 2280 HV Rijswijk  
Tel. (+31-70) 340-2040, Tx. 31 651 epo nl,  
Fax: (+31-70) 340-3016

Authorized officer

Rigondaud, B

# INTERNATIONAL SEARCH REPORT

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PCT/SE 98/01101

**C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT**

Category	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	WO 91 07350 A (EKA NOBEL AB) 30 May 1991  see claims 1-3,6-10 see page 2, line 29 - page 9, line 37	1,4,5, 7-10, 13-18
A	& US 5 368 833 A cited in the application ----	2,3,6,12
A	WO 91 07351 A (EKA NOBEL AB) 30 May 1991 see the whole document & US 5 447 604 A cited in the application ----	1,14-18
A	PATENT ABSTRACTS OF JAPAN vol. 010, no. 123 (C-344), 8 May 1986 & JP 60 251119 A (NISSAN KAGAKU KOGYO KK), 11 December 1985 see abstract ----	1,2,9,10
A	WO 89 06637 A (E. I. DU PONT DE NEMOURS AND COMPANY) 27 July 1989 see claims 1-3,6,15 see page 1, line 4 - line 9 see page 9, line 3 - line 26 see page 14, line 21 - page 15, line 15 & US 5 176 891 A cited in the application -----	1,14-18

# INTERNATIONAL SEARCH REPORT

Information on patent family members

Inte...inal Application No

PCT/SE 98/01101

Patent document cited in search report	Publication date	Patent family member(s)		Publication date
WO 9405596	A	17-03-1994	SE 501214 C AT 138354 T AU 667966 B AU 4988193 A CA 2141551 A CN 1084490 A,B DE 69302823 D DK 656872 T EP 0656872 A ES 2087767 T FI 950622 A JP 2787377 B JP 8502016 T MX 9305272 A NO 950738 A NZ 255615 A RU 2081060 C SE 9202502 A US 5603805 A	12-12-1994 15-06-1996 18-04-1996 29-03-1994 17-03-1994 30-03-1994 27-06-1996 07-10-1996 14-06-1995 16-07-1996 13-02-1995 13-08-1998 05-03-1996 28-02-1994 27-02-1995 27-11-1995 10-06-1997 01-03-1994 18-02-1997
WO 9107350	A	30-05-1991	SE 500387 C AT 107608 T AU 628692 B AU 6733490 A CA 2067506 A,C CN 1115817 A DE 69010210 D DE 69010210 T DK 491879 T EP 0491879 A ES 2055581 T FI 922056 A,B, JP 4505314 T JP 5009368 B KR 9505762 B LT 445 A,B LV 10227 A,B PT 95849 A SE 8903753 A RU 2068809 C	13-06-1994 15-07-1994 17-09-1992 13-06-1991 09-05-1991 31-01-1996 28-07-1994 13-10-1994 07-11-1994 01-07-1992 16-08-1994 06-05-1992 17-09-1992 04-02-1993 30-05-1995 25-10-1994 20-10-1994 13-09-1991 10-05-1991 10-11-1996

# INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No

PCT/SE 98/01101

Patent document cited in search report	Publication date	Patent family member(s)		Publication date
WO 9107350	A	US	5643414 A	01-07-1997
		US	5368833 A	29-11-1994
WO 9107351	A 30-05-1991	SE	500367 C	13-06-1994
		AT	107609 T	15-07-1994
		AU	635365 B	18-03-1993
		AU	6734290 A	13-06-1991
		CA	2067495 A,C	10-05-1991
		DE	69010217 D	28-07-1994
		DE	69010217 T	13-10-1994
		DK	502089 T	07-11-1994
		EP	0502089 A	09-09-1992
		ES	2055580 T	16-08-1994
		FI	922057 A,B,	06-05-1992
		JP	4505315 T	17-09-1992
		SE	8903754 A	10-05-1991
		US	5447604 A	05-09-1995
WO 8906637	A 27-07-1989	AU	3049789 A	11-08-1989
		DE	68914638 D	19-05-1994
		DE	68914638 T	21-07-1994
		EP	0382795 A	22-08-1990
		US	5176891 A	05-01-1993